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AMENDMENTS TO THE CLAIMS

Please amend the claims as follows:

LISTING OF CLAIMS:

1. (Withdrawn) A method of making a nutraceutical composition for the

treatment or prevention of diabetes and/or obesity and syndrome X comprising

admixing a catechin found in green tea and a PPARy ligand to form a nutraceutical

composition.

2. (Withdrawn) A method according to claim 1 wherein the PPARv

ligand is selected from the group consisting of a full agonist, a partial agonist, a

selective PPARy modulator/agonist, and a PPARy dual agonist or panagonist.

3. (Withdrawn) A method according to claim 1 wherein the PPARy ligand

is a thiazolidinedione.

4. (Withdrawn) A method according to claim 1 wherein the PPARv ligand

is a natural PPARy agonist.

5. (Withdrawn) A method according to claim 1 wherein the PPARy ligand

is a PUFA.

6. (Withdrawn) A method according to claim 1 wherein the PPARγ ligand

is ligustilide.

7. (Withdrawn) A method according to claim 1 wherein the PPARy ligand

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is phytanic acid.

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8. (Withdrawn) A method of treating or preventing diabetes and/or obesity

and syndrome X comprising consuming a nutraceutical composition comprising a

catechin found in green tea during administration of a PPARy ligand.

9. (Withdrawn) A method according to claim 8 wherein the nutraceutical

composition is a food or beverage or a supplement composition for a food or beverage.

10. (Withdrawn) A method according to claim 8 wherein the nutraceutical

composition is a pharmaceutical composition.

11. (Withdrawn) A method according to claim 8 wherein the catechin is (-)

epigallocatechin gallate.

12. (Withdrawn) A method for the treatment or prevention of diabetes or

obesity and syndrome X which comprises administering to a subject in need of such

treatment an effective amount of a catechin found in green tea and of a PPARy ligand.

13. (Withdrawn) The method as in claim 12 wherein the catechin is

(-) epigallocatechin gallate.

14. (Previously presented) A composition comprising a catechin found in

green tea, and a peroxisome proliferator-activated receptor gamma (PPARγ) ligand

selected from the group consisting of thiazolidinediones, ligustilide and phytanic acid,

wherein the composition is a pharmaceutical composition.

15. (Original) A composition as in claim 14 wherein the catechin is (-)

epigallocatechin gallate.

16. (Withdrawn): A composition according to claim 14, wherein the

thiazolidinedione is ciglitazone, rosiglitazone or pioglitazone.

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17. (Previously Presented): A composition according to claim 15 wherein

(-) epigallocatechin gallate is present in an amount sufficient to administer to a human

adult a daily dosage of about 10 mg to about 2000 mg.

18. (Canceled).

19. (Withdrawn) A method according to claim 3 wherein the

thiazolidinedione, is selected from the group consisting of ciglitazone, rosiglitazone and

pioglitazone.

20. (Withdrawn) A method according to claim 5 wherein the PUFA is

selected from the group consisting of eicosapentaenoic acid and docosahexaenoic acid.

21. (Previously presented) The composition according to claim 14

wherein the PPARy ligand is ligustilide.

22. (Previously presented) The composition according to claim 14

wherein the PPARy ligand is in a dosage of from about 1 to about 1000 mg.

23. (Currently amended) The composition according to claim 14 wherein

the pharmaceutical composition is a solid unit oral dosage form, the catechin is (-)

epigallocatechin gallate and (-) epigallocatechin gallate is present in an amount of from

about 10 mg to about 2000 mg, and wherein the PPARy ligand is present in an amount

of from about 1 to about 1000 mg.

24. (New) The composition according to claim 14 wherein the

pharmaceutical composition is a solid unit oral dosage form for effecting glucose

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tolerance and preventing body weight gain or adipose tissue weight gain associated with use of a PPARy ligand and the catechin and the PPARy ligand are present in

glucose lowering amounts.

25. (New) A pharmaceutical composition for effecting glucose tolerance

comprising an effective amount for reducing fasted state glucose concentration of a

catechin found in green tea, and an effective amount of a peroxisome proliferator-

activated receptor gamma (PPARy) ligand selected from the group consisting of

thiazolidinediones, liqustilide and phytanic acid, wherein the amounts of the catechin

and the PPARv ligand are such that fasted state glucose is lowered to an extent greater

than that for either the catechin or the PPARy ligand.

26. (New) A pharmaceutical composition for effecting glucose tolerance

comprising an effective amount of a catechin found in green tea, and of a peroxisome

proliferator-activated receptor gamma (PPARy) ligand selected from the group

consisting of thiazolidinediones, ligustilide and phytanic acid, wherein the effective

amount of each of the catechin and the PPARy ligand in combination reduces fasted

state glucose concentration and prevents body weight gain or adipose tissue weight

gain associated with use of a PPARγ ligand.

27. (New) The composition according to claim 23 wherein the PPARγ

ligand is ligustilide.

28. (New) The composition according to claim 24 wherein the PPARy

ligand is ligustilide.

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- (New) The pharmaceutical composition according to claim 25 wherein the PPARy ligand is ligustilide.
- (New) The pharmaceutical composition according to claim 26 wherein the PPARy ligand is ligustilide.
- 31. (New) The pharmaceutical composition according to claim 25 wherein the catechin is (-) epigallocatechin gallate and (-) epigallocatechin gallate is present in an amount of from about 10 mg to about 2000 mg, and wherein the PPARy liquid is present in an amount of from about 1 to about 1000 mg.
- 32. (New) The pharmaceutical composition according to claim 26 wherein the catechin is (-) epigallocatechin gallate and (-) epigallocatechin gallate is present in an amount of from about 10 mg to about 2000 mg, and wherein the PPARy ligand is present in an amount of from about 1 to about 1000 mg.
- 33. (New) The composition according to claim 23 wherein the (-)-epigallocatechin gallate is present in an amount of from 100 mg to 300 mg, and the PPARy ligand is present in an amount of from 8 mg to 100 mg.
- 34. (New) The composition according to claim 23 wherein the (-)-epigallocatechin gallate is present in an amount of about 2000 mg, and the PPARy ligand is present in an amount of about 1000 mg.